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## **AMENDMENTS TO THE CLAIMS**

1. (original): A method to prepare living cells which cells comprise a first fluorescent protein localized to the nucleus and a second fluorescent protein localized to the cytoplasm

wherein said first and second fluorescent proteins emit light of different wavelengths which method comprises

modifying living cells to contain either

- (a) a first expression system for expression of said first fluorescent protein wherein said first fluorescent protein is fused to an amino acid sequence which targets said fusion protein to the nucleus and a second expression system for expression of a second fluorescent protein lacking a nucleus targeting sequence; or
- (b) an expression system that expresses both said first fluorescent protein and second fluorescent protein as described; and

selecting said modified cells for cells that have been stably modified.

- 2. (original): The method of claim 1, wherein in step (a), said cells are first modified with said first expression system and then modified with said second expression system or *vice* versa.
- 3. (original): The method of claim 1, wherein in step (a), the cells are modified with both expression systems simultaneously.
- 4. (original): The method of claim 1, wherein said selecting is by culturing in the presence of an antibiotic or a toxin.
- 5. (currently amended): A colony of living Living cells stably modified to produce a first fluorescent protein fused to an amino acid sequence targeting the nucleus and a second fluorescent protein lacking an amino acid sequence targeting the nucleus;

wherein said first and second fluorescent proteins emit visible light at different wavelengths.

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6. (currently amended): A colony of living Living cells which are modified to contain a first fluorescent protein localized to the nucleus and a second fluorescent protein localized to the cytoplasm wherein said first fluorescent protein and second fluorescent protein are of different colors.

- 7. (currently amended): The <u>colony eells</u> of claim 6, wherein said first fluorescent protein is green and said second fluorescent protein is red.
- 8. (currently amended): The <u>colony eells</u> of claim 5, wherein said amino acid sequence targeting the nucleus is histone H2B.
- 9. (currently amended): A method to determine the cell cycle position of living cells which method comprises assessing the ratio of nuclear area to cytoplasmic area of the cells of the colony of claim 6.
- 10. (original): The method of claim 9, wherein said assessing is performed as a function of time.
- 11. (currently amended): The method of claim 9, wherein said cells of said colony are observed in a living animal.
- 12. (currently amended): A method to determine the effect of an agent on cells, which method comprises

treating a first sample of the <u>colony eells</u> of claim 6 with said agent and observing the effect of said treating on the distribution and/or intensity of radiation emitted from said <u>colony eells</u>.

13. (currently amended): The method of claim 12, which further comprises observing the distribution and/or intensity of radiation emitted from a second sample of said colony said cells that has have not been treated with said agent, and

comparing the observations made on the first sample with those on the second sample.

- 14. (currently amended): The method of claim 12, wherein the distribution and/or intensity are evaluated for being characteristic of dormancy.
- 15. (currently amended): The method of claim 12, wherein said distribution and/or intensity are evaluated for being characteristic of apoptosis.
- 16. (currently amended): The method of claim 12, wherein said distribution and/or intensity are evaluated for being characteristic of stages in the cell cycle.
- 17. (currently amended): A method to determine the location of targeting of an agent <u>as</u> the cytoplasm or nucleus which method comprises treating the <u>colony cells</u> of claim 6 with said agent and observing the distribution and/or intensity of radiation emitted from the cytoplasm and <u>nucleus said cells</u>.
- 18. (original): The method of claim 17, wherein said agent itself is labeled, and said method further comprises directly observing the location of the label.
- 19. (currently amended): A method to determine the proliferation rate of a cell culture which method comprises culturing cells which have been modified to contain a fluorescent protein; and

measuring the fluorescence emitted by said cells as a function of time, whereby the rate of proliferation of said cells is determined, as correlated to the rate of increase of intensity of emitted fluorescence.

- 20. (original): The method of claim 19, wherein said fluorescent protein is a green fluorescent protein (GFP) or a red fluorescent protein (RFP).
  - 21. (original): The method of claim 19, wherein said culture is grown from a single cell.

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22. (currently amended): A method to determine the effect of a test compound on cell proliferation which method comprises

culturing cells in the presence and absence of said test compound, wherein said cells have been modified to contain a fluorescent protein;

measuring the intensity of fluorescence as a function of time in the presence and absence of said compound so as to determine the rate of proliferation in the presence and absence of said compound, as correlated to the rate of increase of intensity of emitted fluorescence; and

comparing the rate of proliferation in the presence and absence of said compound;

wherein a change in the rate of proliferation in the presence as opposed to the absence

wherein a change in the rate of proliferation in the presence as opposed to the absence of said compound identifies said compound as a modulator of cellular proliferation.

- 23. (original): The method of claim 22, wherein said fluorescent protein is a green fluorescent protein (GFP) or a red fluorescent protein (RFP).
- 24. (original): The method of claim 22, wherein said culturing is commenced from a single cell.
- 25. (original): A method to determine the heterogeneity of a tumor, which method comprises culturing a multiplicity of colonies from individual cells or individual groups of cells contained in said tumor; and

determining the rates of proliferation of said cell cultures;

whereby cultures exhibiting different rates of proliferation indicate heterogeneity of said tumor.

26. (currently amended): The method of claim 25, wherein said cells have been modified to contain a fluorescent protein and the rates of proliferation are determined, as correlated to the rate of increase of intensity of emitted fluorescence, by measuring the intensity of emitted fluorescence as a function of time.

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27. (currently amended): The method of <u>claim 26</u> <u>claim 25</u>, wherein said cells have been modified to contain a first fluorescent protein localized to the nucleus and a second fluorescent protein localized to the cytoplasm wherein said first fluorescent protein and second fluorescent protein are of different colors.